# **AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application:

### LISTING OF CLAIMS

1. (Currently amended) A method in a computer system for preventing atypical clinical events related to information identified by DNA testing a person, comprising the steps of:

receiving clinical agent information, the clinical agent information including an identifier of the agent;

accessing a data structure to determine[ing] if a gene variant is known to be associated with one or more atypical events for the clinical agent information; and

inquiring if the person has a <u>stored genetic test result value</u> for the gene <u>variant;</u>, and if not,

accessing hereditary information for the person if the person does not have a genetic test result value for the genetic variant;

utilizing the hereditary information for the person to determine the likelihood the person has the gene variant; and

generating an output including information regarding the likelihood that the person has [a] the gene variant indicative of an atypical event based on the hereditary information.

3. (Currently amended) The method of claim  $\underline{1}[2]$ , wherein the hereditary information includes ethnicity.

### 4. (Canceled)

- 5. (Currently amended) The method of claim 1[4], wherein the accessing of the hereditary information comprises accessing the hereditary information is obtained from an electronic medical record of the person stored within a comprehensive healthcare system.
- 6. (Currently amended) The method of claim 1[2], further comprising the step of initiating a clinical action if a test result value is not available for the person and the information regarding the risks indicates a significant risk that the person carries has the [a] gene variant associated with an atypical event.
- 7. (Original) The method of claim 6, wherein the clinical action is ordering a genetic test.
- 8. (Currently amended) A computer system <u>embodied on one or more computer</u> storage media having computer-executable instructions embodied thereon for preventing atypical clinical events related to information identified by DNA testing a person, comprising:

a receiving component that receives clinical agent information, the clinical agent information including an identifier of the agent;

a first accessing component for accessing a data structure to a determining component that determine[s] if a gene variant is known to be associated with one or more atypical events for the clinical agent information;

an inquiring component that inquires if the person has a <u>stored genetic test</u> result value for the <u>associated gene variant;[,] and</u>

a second accessing component for accessing hereditary information for the person if the person does not have a genetic test result value for the gene variant;

a utilizing component for utilizing the hereditary information for the person to determine the likelihood the person has the gene variant; and

a generating component that generates an output including information regarding the likelihood that the person has [a] the gene variant indicative of an atypical event based on the hereditary information.

## 9. (Canceled)

10. (Currently amended) The computer system of claim 8[9], wherein the hereditary information includes ethnicity.

- 12. (Currently amended) The computer system of claim <u>8</u>[11], wherein <u>the second accessing component accesses</u> the hereditary information <u>is obtained</u> from an electronic medical record of the person stored within a comprehensive healthcare system.
- 13. (Currently amended) The computer system of claim §[9], further comprising an initiating component that initiates a clinical action if a test result value is not available for the person and the information regarding the risks indicates a significant risk that the person has earries [a] the gene variant associated with an atypical event.

14. (Original) The computer system of claim 13, wherein the clinical action is ordering a genetic test.

15. (Currently amended) A computer-readable medium containing instructions for a method for controlling a computer system for preventing atypical clinical events related to information identified by DNA testing a person, the method comprising the steps of:

receiving clinical agent information, the clinical agent information including an identifier of the agent;

accessing a data structure to determine[ing] if a gene variant is known to be associated with one or more atypical events for the clinical agent information; and

inquiring if the person has a <u>stored</u> genetic test result value for the gene <u>variant</u>; and if not,

accessing hereditary information for the person if the person does not have a genetic test result value for the gene variant;

utilizing the hereditary information for the person to determine the likelihood the person has the gene variant; and

generating an output including information regarding the likelihood that the person has [a]the gene variant indicative of an atypical event based upon the hereditary information.

17. (Currently amended) The computer-readable medium of claim <u>15</u>[16], wherein the hereditary information includes ethnicity.

- 19. (Currently amended) The computer-readable medium of claim <u>15</u>[18], wherein the <u>accessing of the</u> hereditary information <u>comprises accessing the hereditary information obtained</u> from an electronic medical record of the person stored within a comprehensive healthcare system.
- 20. (Currently amended) The computer-readable medium of claim <u>15</u>[16], further comprising the step of initiating a clinical action if a test result value is not available for the person and the information regarding the risks indicates a significant risk that the person earries has the [a] gene variant associated with an atypical event.
- 21. (Original) The computer-readable medium of claim 20, wherein the clinical action is ordering a genetic test.